



(19) Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) Publication number:

0 520 690 B1

(12)

EUROPEAN PATENT SPECIFICATION

(45) Date of publication of patent specification: 02.11.95 (51) Int. Cl.⁶: A61K 6/033, A61K 6/00,
A61L 25/00, A61L 27/00

(21) Application number: 92305603.0

(22) Date of filing: 18.06.92

The file contains technical information submitted
after the application was filed and not included in
this specification

(54) Calcium phosphate type hardening material for repairing living hard tissue.

(30) Priority: 26.06.91 JP 154940/91

(43) Date of publication of application:
30.12.92 Bulletin 92/53

(45) Publication of the grant of the patent:
02.11.95 Bulletin 95/44

(84) Designated Contracting States:
DE FR GB NL

(56) References cited:
WO-A-90/00892
DE-A- 3 730 298
US-A- 4 518 430
US-A- 5 149 368

DERWENT JAPANESE PATENTS REPORT 2
May 1990 Derwent Publications Ltd., London,
GB; & JP-A-2 048 479 (TOKUYAMA SODA CO
LTD) 19.02.90

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Description**BACKGROUND OF THE INVENTION**

5 The present invention relates to hardening materials for repairing living hard tissue which are applied to medical cements used for adhesion or fixing for repairing bone defects or gaps, dental pulp capping cements, dental root canal sealing cements, etc.

For medical cements used for adhesion and fixing for repairing bone defects and gaps in orthopaedic surgery field, cyanoacrylate type adhesives and methyl methacrylate polymer-methyl methacrylate monomer type bone cements have been currently employed. These polymers *in vitro*, however, are not able to integrate into hard tissue even after being retained in living body for long periods of time because of their low biocompatibility. On the other hand, for root canal restoration in odontotherapy the typical methods are of filling calcium hydroxide type pastes into root canal or of sealing root apex with zinc oxide eugenol cements together with gutta-percha points. However, there is a problem that the calcium hydroxide type pastes cause irritation because the pH of the neighborhood of the filled becomes too basic. Also, it has been pointed out in various fields that the zinc oxide eugenol cements show high cytotoxicity.

Recently, the granules and the block and porous forms of a calcium phosphate, hydroxyapatite $[Ca_{10}(PO_4)_6(OH)_2]$; it may be called apatite hydroxide or HAP], have been clinically used as materials for repairing bone defects and gaps because of its high biocompatibility, on which many reports have been published.

These materials of hydroxyapatite, however, are leaked easily out from the incised part when filled into bone defects or gaps, because, although hydroxyapatite is nontoxic to living body, it can not integrate into hard tissue even after being retained in living body for long periods of time. In addition, since hydroxyapatite prepared either by dry method or by wet method does not cause hardening reaction, it can not be used as a substitute for the powder component of the above described cements.

The characteristic of cement type materials, that is hardening materials, is of the exceedingly large possibility in applications of the materials because of their hardenability, adhesivity, moldability adaptable to complicate shapes and so on unlike general ceramics (sintered substance).

Because of these reasons, hardening materials using biologically active calcium phosphates have been investigated. Thus, coagulation-hardening type materials, in which the phosphates harden by hydration reaction, and chelation-hardening type materials, in which the phosphates form chelate bonds with organic acids or carboxyl group-containing polymers contained in the liquid component in large amounts, resulting in hardening, have been proposed.

The coagulation-hardening type materials are not necessary to use large amounts of organic acids for hardening, because the materials harden by the phenomena that the calcium phosphates cause hydration in water to coagulate and finally convert into hydroxyapatite or octacalcium phosphate $[Ca_8H_2(PO_4)_6 \cdot 5H_2O]$; it may be called OCP]. Because of this reason, the coagulation-hardening type materials do not cause lowering of pH due to the organic acids.

The biologically active calcium phosphates gradually convert into hydroxyapatite, which is a major component of living hard tissue, for example, in living body or oral cavity and can integrate into living hard tissue. Alpha-tricalcium phosphate $[Ca_3(PO_4)_2]$ (alpha-type); it may be called alpha-TCP), tetracalcium phosphate $[Ca_4(PO_4)_2O]$; it may be called TeCP or 4CP), and so on are exemplified as the phosphates. Various kinds of water-hardening calcium phosphate compositions using these calcium phosphates have been proposed.

45 For the coagulation-hardening type material using tetracalcium phosphate, a coagulation-hardening type dental restorative paste comprising a mixture of tetracalcium phosphate and calcium hydrogen phosphate is proposed. (U.S. Patent No. 4518430).

The conventional coagulation-hardening type materials have problems that they show poor kneading-processability and can not be filled uniformly because their pastes (paste may be called cement mud or kneaded product) do not spread uniformly. In general, hardening materials are used by suitably changing the ratio of their powder to liquid components in order to control setting time. If the amount of the powder component were increased in order to shorten the setting time, the kneading-processability and filling properties of the coagulation-hardening type materials would become further poor.

Because of the reason, the hardening materials which show desired effect only when the powder and liquid components are kneaded in a range of extremely limited ratio are not useful practically.

It was found by the inventors' research that since the pH of the neighborhood of the filled pastes of the conventional coagulation-hardening type materials lay in acidic or highly basic regions during hardening, the materials irritated living body.

According to the present invention there is provided a hardenable, biocompatible composition for repairing living hard tissue comprising a powder component containing $\text{Ca}_4\text{P}_2\text{O}_9$ and at least one of CaHPO_4 and $\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$ at a molar ratio of Ca:P of 1.16-1.95; and a liquid component containing a secondary phosphate ion in a concentration of 1mM to 2M as a secondary phosphate and an organic acid ion in a concentration of 1mM to 2M as an organic acid or as a total of an organic acid and its salt, wherein a weight ratio of the powder component : the liquid component is 0.5-4.0.

There is no special restriction in the production method of the present invention, and those prepared by any type of the production method may be used. For example, CaCO_3 , CaO , $\text{Ca}(\text{OH})_2$, etc as Ca sources, P_2O_5 , H_3PO_4 , $\text{NH}_4\text{H}_2\text{PO}_4$, $(\text{NH}_4)_2\text{HPO}_4$, etc as P sources, and $\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$, CaHPO_4 , $\text{Ca}(\text{H}_2\text{PO}_4)_2$, $\text{Ca}_2\text{P}_2\text{O}_7$ etc as both sources may be used as materials. Combinations of these materials may be used. The dry preparation process is preferred in which $\text{Ca}_2\text{P}_2\text{O}_7$ prepared by sintering $\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$ is mixed with CaO prepared by sintering CaCO_3 in a molar ratio of 1:2 followed by sintering. The tetracalcium phosphate powder prepared by the dry method is preferred for the powder material of cements.

For calcium hydrogen phosphate used in the present invention, the both of its dihydrate ($\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$; it may be called DCPD and its mineral name is bragite) and anhydrous (CaHPO_4 ; it may be called DCPA and its mineral name is monetite) may be employed. Though calcium hydrogen phosphate also has no special restriction in its production method, the grade in the Japanese Pharmacopoeia is preferred.

In the present invention, not only two of tetracalcium phosphate and calcium hydrogen phosphate but also mixtures of three kinds or more of tetracalcium phosphate, calcium hydrogen phosphate, and the other calcium phosphate(s) may be used as the powder component. For example, hydroxyapatite, apatite carbonate, alpha-, alpha'-, beta-, and gamma-tricalcium phosphates, octacalcium phosphate, and the like are given as the other calcium phosphates.

The ratio of tetracalcium phosphate to calcium hydrogen phosphate is preferably in a range of 1.28 to 1.91. Out of this range may cause too long setting time of cement.

The ratio of the total weight of tetracalcium phosphate and calcium hydrogen phosphate to the other calcium phosphate is preferably 0.5 or more in the molar ratio of the total weight to the other calcium phosphate, more preferably 1 or more. Out of this range also may cause too long setting time of cement.

In the present invention, although there is no special restriction in the particle size of the powder component, in order to improve the kneading-processability of the hardening materials as easy as possible and to quicken the setting rate of the hardening materials as rapid as possible, the mean particle size is preferably 50 μm or less, more preferably in a range of 0.1 to 20 μm .

The liquid component used for kneading the above powder component is an aqueous solution containing secondary phosphate and organic acid ions. One or plural kinds selected from water-soluble secondary phosphates such as disodium, dipotassium, and diammonium hydrogen phosphates is given as the secondary phosphate ion sources. As to the organic acid ion sources, one or plural kinds selected from organic acids and carboxyl group-containing polymers such as citric acid, malic acid, malonic acid, maleic acid, fumaric acid, fumaleic acid, lactic acid, acetic acid, oxalacetic acid, isocitric acid, aconitic acid, succinic acid, acrylic acid homopolymer, acrylic acid-itaconic acid copolymer, etc., and their salts may be used. The organic acids produced in the citric acid cycle of the metabolic cycle *in vivo* or their salts are preferred.

The concentration of the secondary phosphate ion in the liquid component is preferably 1 mM to 2 M as secondary phosphate, more preferably 20 to 500 mM, and that of the organic acid ion in the liquid component is preferably 1 mM to 2 M as the organic acid or as the total of organic acid and its salt, more preferably 1 to 500 mM. The lower and higher concentrations than these ranges may cause too slow and too rapid setting rate of the resulting cement, respectively.

The secondary phosphate and organic acid and/or its salt are used preferably in a mixture ratio of (molar concentration of secondary phosphate)/(total molar concentration of organic acid and/or its salt) of 0.02 to 142, more preferably 0.04 to 71. Out of this range in the molar ratio may cause no buffer capacity of the liquid component. If the molar ratio were lower than the above range, it would be impossible to maintain the pH of the neighborhood of the paste in weekly basic regions, and the higher molar ratio may cause poor kneading-processability.

The forementioned liquid component is, for example, a buffer solution containing at least one kind of disodium hydrogen phosphate and dipotassium hydrogen phosphate and containing at least one kind of citric acid and malic acid.

In the hardening materials in the present invention, for example, the above powder and liquid components are kneaded preferably in a (powder weight)/(liquid weight) ratio of 1 to 2.4. If the powder were more than this range, it would be impossible to knead the whole quantity of the powder with the liquid, and

too much liquid may cause too much flowability of the paste and inability in molding and hardening.

The hardening materials in the present invention may optionally contain a thickner to improve moldability and uniformity in filling of their pastes. As to the thickner, for example, one or plural kinds selected from water-soluble polymers such as carboxymethyl cellulose, carboxymethyl cellulose sodium, hydroxypropyl cellulose, polyvinyl alcohol, polyglutamic acid and its salts, and polyaspartic acid and its salts, etc. are proposed. At least one selected from carboxymethyl cellulose sodium, hydroxypropyl cellulose, polyglutamic acid, and polyglutamic acid salts is preferred because of their high solubility in water and moderate viscosity. The thickner may be mixed with the powder or liquid component or the paste during kneading. The preferred ratio in use depends on the type of thickner used. For example, polyglutamic acid and its salts are used preferably at 0.5 to 15 % by weight to the total weight of the liquid component, more preferably at 1 to 10 % by weight, and the other thickeners are used preferably at 0.5 to 10 % by weight, more preferably at 0.5 to 5 % by weight.

The hardening materials in the present invention may contain an X-ray contrast medium optionally. The use of the medium is capable of monitoring upon filling the pastes of the hardening materials and searching the change after filling. One or plural kinds selected from barium sulfate, bismuth(III) carbonate (bismuth oxycarbonate), iodoform, barium apatite, barium titanate, etc. is given for the X-ray contrast medium. It may be mixed with the powder or liquid component or the paste during kneading. For example, its amount for use at 5 to 30 % by weight to the total weight of the powder component is preferred.

The hardening materials in the present invention may contain starch, glucosaminoglycane, alginic acid, chitin, chitosan, heparin, etc. as polysaccharides other than cellulose, collagen, gelatin, and their derivatives as proteins, and antirheumatic agents, antiphlogistics, antibiotics, anti-cancer agents, bone-inducing factors, tannic acid and its derivatives such as catechin, retinoic acid and its derivatives, Depentaplegic-Vg-related family (which may be called DVR gene family), TGF- β 1, TGF- β 2, TGF- β family, etc. in addition to the above essential and optional components.

The hardening materials in the present invention are used in the same manner as for the conventional hardening materials employed in the medical and dental fields. For example, the above essential and optional components are kneaded to form a paste. Since the paste hardens, for example, in 5 to 30 minutes to be unable to deform, the paste should be filled into defect according to the usual method before completion of hardening. The filled paste hardens under living body or oral cavity environment and converts completely into hydroxyapatite, for example, in 1 day, and the hardened substance is partially displaced into new bone and finally integrates into living hard tissue.

The hardening materials in the present invention are used, for example, as medical cements for adhesion or fixing, dental pulp capping cements, dental root canal sealing cements and so on.

When used in the above applications, in the comparison with the conventional hardening materials, the hardening materials of the present invention are excellent in the following points: superior kneading-processability to that of the conventional hardening materials, good filling properties of their pastes, controllable setting time in the region longer than few minutes, stable pH of the neighborhood of their pastes in a weakly basic region from the initial stage, conversion of their pastes into hydroxyapatite in living body or oral cavity in short periods of time, promotion of forming living hard tissue such as new bone in the neighborhood of the filled, and finally integrating into living hard tissue to such an extent that the face (interface) where the paste comes in contact with the hard tissue can not be perceived because the paste surface is replaced by the new bone.

The hardening materials in the present invention convert into hydroxyapatite in short periods of time, because the powder component contains at least tetracalcium phosphate and calcium hydrogen phosphate.

Because the liquid component contains secondary phosphate and organic acid ions, the setting time of the materials is reduced and the pH lies in a weakly basic region (for example pH 8.0 to 9.0) from the initial stage of hardening of the materials and is stabilized in the region.

The hardening materials in the present invention show excellent kneading-processability and uniform filling property of their pastes and can be adjusted at moderate setting time. In addition, the pH of the neighborhood of the filled paste part immediately after filling into living body lies in a weakly basic region, and the materials are capable of integrating into living hard tissue.

When the hardening materials contain the thickner as well, they have an advantage of being further improved in the kneading-processability.

When the hardening materials contain the X-ray contrast medium as well, the conversion of them into living hard tissue can be monitored by radiography.

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BRIEF DESCRIPTION OF THE DRAWING

Fig. 1 shows X-ray diffraction patterns of the hardened products from the hardening materials in Example 1 and Reference Examples 4 and 5.

5 Fig. 2 shows time-course of changes in pH of the neighborhood of the pastes from the hardening materials in Example 1 and Reference Examples 4 and 5 from the start of kneading.

Fig. 3 shows time-course of changes in pH of the neighborhood of the pastes from the hardening materials in Examples 2 and 5 and Reference Examples 6-8 from the start of kneading.

10 Fig. 4 shows time-course of changes in pH of the neighborhood of the pastes from the hardening materials using different liquid components (Example 31 and Reference Examples 21-25) from the start of kneading.

Fig. 5 shows time-course of changes in pH of the neighborhood of the pastes from the hardening material in Example 31 and the materials using different powder components from the start of kneading.

15 Fig. 6 shows time-course of changes in pH of the neighborhood of the pastes from the hardening material in Example 32 and the materials using different powder components from the start of kneading.

Fig. 7 shows time-course of changes in pH of the neighborhood of the pastes from the hardening material in Reference Example 21 and the materials using different powder components from the start of kneading.

20 Fig. 8 shows time-course of changes in pH of the neighborhood of the pastes from the hardening material in Reference Example 23 and the materials using different powder components from the start of kneading.

Fig. 9 shows time-course of changes in pH of the neighborhood of the pastes from the hardening material in Reference Example 24 and the materials using different powder components from the start of kneading.

25 Fig. 10 shows time-course of changes in pH of the neighborhood of the pastes from the hardening material in Reference Example 25 and the materials using different powder components from the start of kneading.

Fig. 11 shows time-course of changes in pH of the neighborhood of the pastes from the hardening material in Reference Example 26 and the materials using different powder components from the start of 30 kneading.

Fig. 12 shows time-course of changes in pH of the neighborhood of the pastes from the hardening material in Reference Example 27 and the materials using different powder components from the start of kneading.

35 DESCRIPTION OF THE PREFERRED EMBODIMENTS

Hereinafter, concrete examples and reference examples of the present invention are shown, but the invention is not limited to the below-described examples.

40 (Method of Measurement)

1. Mean particle size

The particles obtained were dispersed in isopropyl alcohol followed by measuring the particle size by 45 means of a particle size distribution meter (CAPA-700; made by Horiba Seisakusho), in which the principle of the method is centrifugal sedimentation.

2. Structure

50 The structure of tetracalcium phosphate produced and the reaction products obtained by kneading and hardening the hardening materials was identified by using a X-ray powder diffraction system (MXP³; made by Max Science Co.).

3. Setting time of hardening material

55 The measurement is based on the method of measuring setting time of dental root canal filling materials in the International Organization for Standardization Standard. Thus, pastes obtained by kneading hardening materials for 1 minute were filled in a ring mold with a diameter of 10 mm and a height of 2 mm

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and, after 2 minutes from the start of kneading, measured by using a Gilmore-type needle with a mass of 100 g and a diameter 2 mm under a condition of 37°C and a relative humidity of not less than 95 %. The time until indentations cease to be visible was measured as setting time. The measurement was done up to 2 hours, and the time exceeding 2 hours was expressed as 'not hardening'.

5

4. pH of neighborhood of paste

The pH was measured by using a compact pH meter (CARDY; made by Horiba Seisakusho). Briefly, a water-absorbing paper was placed on the sensor of the pH meter, 2 or 3 drops of distilled water were 10 dropped thereon to wet the paper, and pastes obtained by kneading hardening material for 1 minute were placed on the wetted paper. The pH values measured at 3, 5, 10, 15, and 20 minutes after the start of kneading at room temperature were evaluated as the pH of the neighborhood of the pastes.

5. Kneading processability

15

Materials from which the pastes adhered to the kneading spatula during kneading were judged as "good kneading processability". The other judgements were based on feeling in touch.

6. Integration into living hard tissue

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Pastes were filled into root canal of the supramandibular first molar of SD rats as direct capping agents, the rats were bred and sacrificed, and sections were prepared in accordance with the usual method. The evaluation was pathologic-histologically done with the guidance of calcification in the root canal pulp.

25 (Example 1 for preparation of powder component)

Calcium pyrophosphate ($\text{Ca}_2\text{P}_2\text{O}_7$) prepared by sintering calcium hydrogen phosphate dihydrate ($\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$; made by Hoeiyakkou Co. Ltd.; Grade in the Japanese Pharmacopoeia) at 1,100°C was mixed with calcium oxide prepared by sintering calcium carbonate (CaCO_3 ; made by Ebisu Yakuhin Kogyo Co. Ltd.; Grade in the Japanese Pharmacopoeia) at 1,100°C in a molar ratio of 1:2 and then sintered at 30 1,400°C. The resulting product was pulverized by using a ball mill and then classified to recover particles with a particle size of not larger than 32 μm . Thus obtained powder was confirmed as pure tetracalcium phosphate by the identification using the X-ray powder diffraction system, in which all the peaks obtained were equal to those of the tetracalcium phosphate in the JSPD Card No. 25-1137.

35 The tetracalcium phosphate (mean particle size 9.2 μm) and calcium hydrogen phosphate (mean particle size 5.8 μm ; the above described calcium hydrogen phosphate dihydrate) which had been classified to a particle size of not larger than 32 μm were mixed in the various molar ratios shown in Tables 2-4 to give powder components (powder material sample).

40 (Preparation of liquid component)

Aqueous solutions were prepared using the constituents at concentrations shown in Table 1 and used in the following examples and reference examples.

45

TABLE 1

50

Na ₂ HPO ₄ -CITRIC ACID BUFFER SOLUTION				
No.	I	II	III	IV
M/5 Na ₂ HPO ₄ (ml)	10.30	12.63	16.47	19.45
M/10 citric acid (ml)	9.70	7.37	3.53	0.55
pH	5.0	6.0	7.0	8.0

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EXAMPLES 1-9, 12-15 and REFERENCE EXAMPLES 1-3

Hardening materials were prepared using the powder and liquid components shown in Tables 2-4 at the powder to liquid ratios (powder weight/liquid weight) shown in Tables 2-4.

5

REFERENCE EXAMPLE 4

A hardening material was prepared in the same manner as Example 1 except for using distilled water as the liquid component.

10

REFERENCE EXAMPLE 5

A hardening material was prepared in the same manner as Example 1 except for using 20 mM orthophosphoric acid as the liquid component.

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REFERENCE EXAMPLE 6

A hardening material was prepared in the same manner as Example 2 except for using, as the liquid component, a glycine-NaCl-HCl buffer solution (pH 1.42) of the following composition:

20

0.1 M glycine + 0.1 M NaCl	3.0 mL
0.1 N HCl	7.0 mL

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REFERENCE EXAMPLE 7

A hardening material was prepared in the same manner as Example 2 except for using, as the liquid component, the disodium citrate-HCl buffer solution (pH 4.89) of the following composition:

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0.1 M disodium citrate	9.5 mL
0.1 N HCl	0.5 mL

35

REFERENCE EXAMPLE 8

A hardening material was prepared in the same manner as Example 2 except for using, as the liquid component, a glycine-NaCl-NaOH buffer solution (pH 11.25) of the following composition:

40

0.1 M glycine + 0.1 M NaCl	5.0 mL
0.1 N NaOH	5.0 mL

45

EXAMPLE 10

A hardening material was prepared using the powder component prepared as the above-described method using the molar ratio shown in Table 3 and the disodium hydrogen phosphate-citric acid buffer solution (pH 5.0) containing 2.5 % by weight of carboxymethyl cellulose sodium (shown as CMCNa in the Table) in a powder to liquid weight ratio of 1.4.

REFERENCE EXAMPLE 9

55 A hardening material was prepared in the same manner as example 10 except for using distilled water containing 2.5 % by weight of carboxymethyl cellulose as the liquid component and a powder to liquid weight ratio of 1.0.

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EXAMPLE 11

A hardening material was prepared in the same manner as example 10 except using a X-ray contrast medium (compound name; iodoform) at 30 % by weight to the total weight of the powder component.

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REFERENCE EXAMPLE 10

A hardening material was prepared using calcium oxide as the powder component and physiological saline as the liquid component.

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EXAMPLES 16-30 and REFERENCE EXAMPLES 11-20

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The hardening materials prepared in Examples 1-15 and Reference Examples 1-10 were kneaded to examine kneading-processability, setting time, and pH of the neighborhood of the pastes in accordance with the above described methods. The results are shown in Tables 2-4.

In the tables, 4CP, DCPD, DSP, DPP, DSC, HAp, CA, and MA represent tetracalcium phosphate, calcium hydrogen phosphate dihydrate, disodium hydrogen phosphate 12 hydrate, dipotassium hydrogen phosphate anhydrous, disodium citrate, hydroxyapatite, citric acid, and malic acid, respectively.

20

TABLE 2

25

		EXAMPLE								
		1	2	3	4	5	6	7	8	9
Powder Component	4CP:DCPD Molar Ratio	1:1	1:1	1:1	1:1	1:1	1:3	3:1	1:5	5:1
Liquid Component	DSP-CA Buffer Soln. (No.)	I	I	II	III	IV	I	I	I	I
Powder/Liquid Wt. Ratio		1.0	1.6	1.6	1.6	1.6	1.6	1.6	1.6	1.6
Kneading-Processability		good	good	good	good	good	good	good	good	good
Setting Time (min.)		105	45	18	8	7	38	45	36	41
pH of Neighborhood of past	After 3 min.	8.03	8.61	8.59	9.08	8.77	8.57	7.85	8.76	7.70
	After 20 min.	8.53	8.80	8.81	8.98	8.63	8.66	8.51	8.86	8.31

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TABLE 3

		EXAMPLE					
		10	11	12	13	14	15
5 Powder Component (Molar Ratio)	4CP	1	1	2	1	1	1
	DCPD	1	1	2	1	1	1
	HAp	---	---	1	---	---	---
	CHI, (wt.%)	---	30	---	---	---	---
10 Liquid Component (Wt. %)	DSP	I	I	I	10	---	10
	DPP	in	in	in	---	10	---
	CA	Tab.1	Tab.1	Tab.1	5	5	---
	MA				---	---	5
Other Component		CMCNa	CMCNa	CMCNa	CMCNa	CMCNa	CMCNa
Powder/Liquid Wt. Ratio		1.4	1.4	1.2	1.2	1.2	1.2
Kneading-Processability		good	good	good	good	good	average
Setting Time (min.)		42	53	46	9	10	14
20 pH of Neighbor- hood of paste	After 3 min.	8.40	8.51	8.26	7.61	7.81	7.87
	After 20 min.	8.76	8.78	8.62	8.07	8.20	7.94

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TABLE 4
REFERENCE EXAMPLE

		1	2	3	4	5	6	7	8	9	10
Powder Component	4CP:DCPD Molar Ratio	1:0	1:1	1:1	1:1	1:1	1:1	1:1	1:1	1:1	Ca (OH) ₂
Liquid Component	I. 0.2M in DSP CA	0.1M	D.W.	20mM H ₃ PO ₄	DSC-HCl	Gly-NaCl-NaOH	D.W.	D.W.	D.W.	P.S.	
Other Component	---	---	---	Soln. Soln.	HCl	NaOH	---	---	---	CMCNa	---
Powder/Liquid Wt.-Ratio	1.6	1.0	1.0	1.0	1.0	1.6	1.6	1.6	1.0	1.4	
Kneading Processability	poor	poor	poor	poor	poor	age	age	age	age	age	average
Setting Time (min.)	n.h.	28	n.h.	n.h.	n.h.	44	53	29	---	lump formation	formation
pH of Neighborhood of Paste	After 3 min.	6.76	9.11	5.70	12.19	11.48	12.31	12.53	12.69	---	n.h.
	After 20 min.										12.89

D.W.: distilled water

Gly-NaCl-HCl: Glycine-NaCl-HCl buffer solution

DSC-HCl: disodium citrate-HCl buffer solution

Gly-NaCl-NaOH: Glycine-NaCl-NaOH buffer solution

P.S.: Physiological saline

n.h.: not hardened

As can be seen in Tables 2-4, the hardening materials in these examples showed good kneading-processability, the setting time was controllable at 7 minutes or more, and the pH of the neighborhood of their pastes lay in weakly basic regions at the initial stage of hardening from the start of kneading.

In Figure 1, the results of the identification using the X-ray powder diffraction system for the hardened products prepared by allowing the pastes from the hardening materials in Example 1 and Reference Examples 4 and 5 to stand for 16 hours in an atmosphere of 37°C and a relative humidity of more than 95 % are shown.

In Figure 2, the time-course of the changes in pH of the neighborhood of the pastes from the hardening materials in Example 1 and Reference Examples 4 and 5 from the start of kneading is shown.

In Figure 3, the time-course of those in Example 2 and 5 and Reference Examples 6-8 is shown.

As can be seen in curve 1 shown in Figure 2, for the hardening material (Example 1), which was kneaded with the disodium hydrogen phosphate-citric acid buffer solution (pH 5.0), the pH of the neighborhood of the paste was stabilized at a weakly basic (pH about 8.5) at 20 minutes after the start of

kneading. Also, for this hardening material the powder component converted into poorly crystalline hydroxyapatite at 16 hours after the start of kneading, as shown in pattern 11 in Figure 1. In contrast, for the hardening materials using distilled water (Reference Example 4; curve 2 in Figure 2) and 20 mM orthophosphoric acid solution (Reference Example 5; curve 3 in Figure 2), respectively, the pH of the neighborhood of these pastes was both stabilized at a highly basic (pH about 12.5) at 20 minutes after the start of kneading. In addition no change in these powder components was observed after 16 hours [patterns 12 (Reference Example 4) and 13 (Reference Example 5) in Figure 1]. The pattern 14 in Figure 1 shows the result of the powder component used in Example 1.

For the hardening materials (Examples 2-5) which were kneaded with the disodium hydrogen phosphate-citric acid buffer solutions (pH 5.0-8.0), when the pH of the buffer solution is 5.0, the setting time was increased and, when the pH of that is 8.0, the setting time was reduced. Thus, it was shown that higher basic in pH of the liquid tended to shorter setting time. In addition the pH of the neighborhood of these pastes was stabilized at a weakly basic region (pH 8.0-9.0) at 20 minutes after kneading in all the pH of the solutions [curves 4 (Example 2) and 5 (Example 5) in Figure 3]. In contrast, when using the glycine-NaCl-HCl, glycine-NaCl-NaOH, and disodium citrate-HCl buffer solutions, respectively, the pastes hardened in 1 hour, but the pH of the neighborhood of these pastes was stabilized at a highly basic region (pH 12.5-13.0) in all the buffer solutions [curves 6 (Reference Example 6), 7 (Reference Example 7), and 8 (Reference Example 8) in Figure 3].

When distilled water containing 2.5 % by weight of carboxymethyl cellulose sodium was used as the liquid component, the hardening material formed lumps during kneading and the processability was poor (Reference Example 9).

Subsequently, the integration into living hard tissue for the hardening materials in Examples 2, 10, 11, 13, and 14 and Reference Example 10 and the identification of the hardened products by X-ray powder diffraction for those in Examples 2, 10, 11, 13, and 14 and Reference Examples 4 and 5 were investigated.

The integration into living hard tissue was examined in such a manner that hardening materials (pastes) were filled into root canal of SD rats as direct capping agents and the pulp reaction was pathologic-histologically evaluated. Thus, it was examined whether there were lesions or not in the root apex and calcification occurred or not (when calcification was observed, the time when it occurred and the place in the root canal where it occurred were also examined). These results were compared with that of the hardening material using calcium hydroxide as the powder component (Reference Example 10), which has at present been used as a pulp capping agent in the dental field.

Thus, a pulp chamber of the supramandibular first molar of SD rats was opened by using a #1/2 round bar (a dental rotating cutting machine which is one of cutting tool for electric engine) (to make a hole in the molar so as to penetrate to the pulp cavity). The pulp cavity from which the corona dentis pulp was mechanically removed was filled with the material and sealed with a glass-ionomer cement (ones of dental cements used for adhesion of metallic restorative dental materials, pulp capping, filling anterior teeth, and so on). The molar filled with a sterilized type and retained opened pulp cavity until completion of the experiment was used as a control. The rats were sacrificed 1, 2, 3, 4, and 5 weeks after bred, the mandibulae were taken out, demineralized, and paraffin-embedded as the usual method, and continuous sections with a thickness of approximately 5 µm were made, hematoxylin orange-dyed, and then examined pathologic-histologically.

The identification of the hardened products by X-ray powder diffraction was performed in such a manner that hardening materials were kneaded and then allowed to stand for 16 hours in an atmosphere of 37°C and a relative humidity of not less than 95%, and the resulting hardened products were subjected to the X-ray powder diffraction system to examine whether the change of the powder components occurred or not.

As the results the pulp necrosed with the elapse of time and the formation of apical lesions was observed after 4 weeks for the control. In contrast, for the hardening materials in Examples 2, 10, 11, 13, and 14, marked addition of hard tissue to the root canal wall was found after 3 weeks, and the whole of the root canal was sealed by hard tissue after 5 weeks. Inflammation and necrosis of the pulp tissue were also not recognized during the experiment in these examples. For the hardening material in Reference Example 10 which comprises calcium hydroxide kneaded with physiological saline, although the material was calcified at the corona dentis side of the upper area of the root canal to form dentin bridge (a scientific name of a hard tissue having dentin-like structure) after 2-4 weeks, congestion was found at the middle area of the root canal after 3 weeks. As the results of X-ray powder diffraction, all the hardened products in these examples converted into hydroxyapatite at 16 hours after kneading. The above-described dentin bridge formed by calcium hydroxide has a structure which causes difficulty in mechanical treatment of root canal that is necessary frequently and subsequently and, therefore, is not always to be a desired pulp reaction. In

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contrast, since the whole of root canal was sealed by hard tissue as shown above when the hardening materials for repairing living hard tissue in the present invention were used, the materials are useful as medial cements, dental pulp capping cements, dental canal sealing cements and the like for adhesion or fixing which utilize the characteristics of inducing the formation of hard tissue.

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(Example 2 for preparation of powder component)

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A powder component is prepared in the same manner as the example 1 for preparation of powder component except for using a 1:1 molar ratio mixture of tetracalcium phosphate having a mean particle size of 15.8 μm and calcium hydrogen phosphate dihydrate having a mean particle size of 5.9 μm and used in the following examples and reference examples. (Hereinafter, this powder component is referred to as "TeDCPD".)

EXAMPLES 31-35

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Hardening materials in the present invention were prepared using the combinations of TeDCPD and the buffer solutions shown in Table 5.

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TABLE 5

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No.	Constituent and pH Value
Na ₂ HPO ₄ -citric acid type buffer solution	
1	0.2M Na ₂ HPO ₄ 10.30 ml + 0.1M citric acid 9.70 ml (pH 5)
2	0.4M Na ₂ HPO ₄ 10.30 ml + 0.2M citric acid 9.70 ml (pH 5)
Na ₂ HPO ₄ -malic acid type buffer solution	
3	0.2M Na ₂ HPO ₄ 10.30 ml + 0.1M malic acid 9.70 ml (pH 5.8)
4	0.4M Na ₂ HPO ₄ 10.30 ml + 0.2M malic acid 9.70 ml (pH 5.8)
K ₂ HPO ₄ -citric acid type buffer solution	
5	0.4M K ₂ HPO ₄ 10.30 ml + 0.2M citric acid 9.70 ml (pH 5)

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REFERENCE EXAMPLES 21-27

Hardening materials were prepared using the combinations of TeDCPD and the buffer solutions shown in Tables 6 and 7.

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The materials in Examples 31-35 and Reference Examples 21-27 were kneaded at the powder to liquid ratios (powder weight/liquid weight) shown in Tables 6 and 7 and examined kneading-processability, setting time, and pH of the neighborhood of the pastes (at 5 and 20 minutes after the start of kneading) as above described. The results are shown in Tables 6 and 7, in which excellent in kneading-processability means occurrence of viscosity during kneading.

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TABLE 6

Liquid Component	EXAMPLE 31		EXAMPLE 32	
	(1) in Tab. 5	(2) in Tab. 5	(2) in Tab. 5	(2) in Tab. 5
Powder/Liquid Wt. Ratio	1.6	1.0	1.6	2.0
Kneading-Processability	excellent	good	excellent	average
Setting Time (min.)	40	24	20	11
pH of Neighbor- hood of Paste	After 5 min. After 20 min.	8.59 8.69	8.49 8.60	8.62 8.68
				8.43 8.56
				8.46 8.57

Liquid Component	EXAMPLE 33		EXAMPLE 34		EXAMPLE 35		REFERENCE	
	(3) in Tab.5	(4) in Tab.5	(5) in Tab.5	(5) in distilled water	EXAMPLE 21	EXAMPLE 22	REFERENCE	REFERENCE
Powder/Liquid Wt. Ratio	1.6	1.6	1.6	1.0	1.0	1.0	1.6	1.6
Kneading-Processability	excellent	excellent	excellent	poor	poor	poor	poor	poor
Setting Time (min.)	10	7	14	n.h.	26	24		
pH of Neighbor- hood of Paste	After 5 min. After 20 min.	8.47 8.56	8.34 8.39	8.51 8.62	12.35 12.48	12.50 12.69	12.26 12.40	

n.h.; not hardened

TABLE 7

		REFERENCE EXAMPLE				
		23	24	25	26	27
5	Liquid Component	20mM		0.2M	0.4M	0.1M
		H_3PO_4		DSP	DSP	CA
10	Powder/Liquid Wt. Ratio	1.0	1.6	1.6	1.6	1.6
Kneading Processability	poor	average	average	average	good	good
Setting Time (min.)	n.h.	19	6	8	n.h.	n.h.
15	pH of Neighborhood of Paste	After 5 min.	8.31	12.29	9.00	9.23
		After 20 min.	8.34	12.17	9.04	8.73
n.h.; not hardened						

The changes in pH of the neighborhood of the pastes from the materials using a powder to liquid ratio of 1.6 in Example 31 and Reference Examples 21-25 are shown in Figure 4. The curves 31, 21, 22, 23, 24, and 25 in Figure 4 correspond to Example 31 and Reference Examples 21, 22, 23, 24, and 25, respectively.

The followings are clear from the results shown in Tables 6 and 7 and Figure 4:

The hardening materials in Examples 31-35 produced viscosity during kneading and hardened in 1 hour while the pH of the neighborhood of the pastes was maintained at a weakly basic region throughout after the start of kneading. The occurrence of viscosity during kneading smoothes kneading, and hence the operation of kneading becomes easy. Furthermore, since the paste having viscosity shows good spreadability, the paste can be filled uniformly and easily.

For the both of the materials in Reference Example 21 which were kneaded with distilled water, the kneading-processability was poor and the pH of the neighborhood of the pastes lay in a highly basic region even on changing the powder to liquid ratio.

For the material in Reference Example 22 which was kneaded with physiological saline, the kneading-processability was poor and the pH of the neighborhood of the paste lay in a highly basic region until completion of hardening from the start of kneading.

The materials in Reference Example 23 which were kneaded with orthophosphoric acid did not harden in a powder to liquid ratio of 1.0, and the pH of the neighborhood of the paste in a ratio of 1.6, which hardened, lay in a highly basic region until completion of hardening from the start of kneading. The material in Reference Example 23 which was kneaded in a powder to liquid ratio of 1.0 was of the same condition as reference Example 5, and hence the same result would be expected. The pH of the neighborhood of the paste, however, was largely different from each other. It is assumed that the use of the powder component having different particle size causes the different result.

When the pastes showing highly basic in the pH of the neighborhood of the pastes such as in Reference Examples 21-23 are filled into a hemorrhagic diseased part, they cause hemolysis and congestion, and hardening of them are inhibited. In addition, the pastes flow out from the diseased part with the bled blood because they have no hemostatic function.

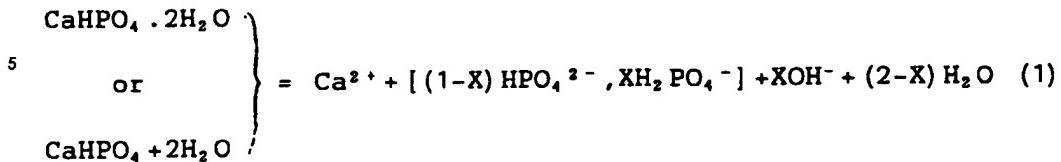
For the materials in Reference Examples 24 and 25 which were kneaded with the disodium hydrogen phosphate solutions, the smoothness on kneading was poor and the pH of the neighborhood of the pastes was slightly higher basic at the initial stage in comparison with those in Examples 31-35.

The materials in Reference Examples 26 and 27 caused neither coagulation-hardening nor chelation-hardening entirely because the concentration of the citric acid solution was low.

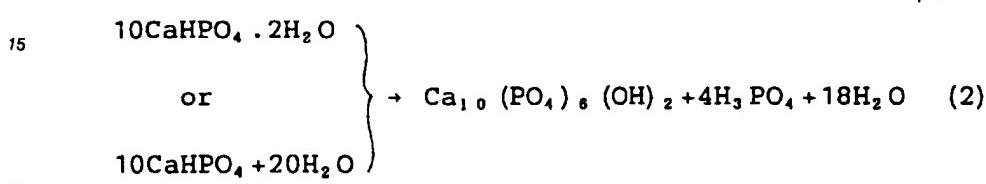
The materials in Examples 31 and 32 and Reference Examples 21 and 23-27 were kneaded in a powder to liquid ratio of 1.0 and examined the time-course (at 3, 5, 10, 15, 20, 25, 30, 40, 50, and 60 minutes after the start of kneading) of the changes in the pH of the neighborhood of these pastes. The results are shown in Figures 5-12. In each figure, the changes in the pH of the neighborhood of the pastes prepared by kneading each of the tetracalcium phosphate and calcium hydrogen phosphate, which were used in the example 2 for preparation of powder component, with the same liquid component as used for kneading TeDCPD were similarly examined and shown. In these figures, curves TeCP and DCPD correspond to the pH changes on kneading the tetracalcium phosphate and calcium hydrogen phosphate, respectively.

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Calcium hydrogen phosphate dissolves in distilled water and attains equilibrium as the formula 1 (see Figure 7):

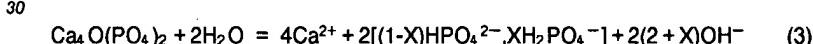


10 The equilibrated solution is a saturated aqueous calcium hydrogen phosphate solution and its calculated pH value is 8.8. Calcium hydrogen phosphate hydrates in the equilibrated solution as shown in the formula 2 to convert into hydroxyapatite:



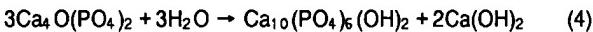
Since this equilibrated solution is supersaturated with respect to hydroxyapatite, the formed hydroxyapatite is deposited. When the conversion shown in the formula 2 proceeds, the pH of the solution rapidly lowers with the deposition of hydroxyapatite and attains equilibrium at the near crossing (approximately pH 4.5) of the solubility curves of calcium hydrogen phosphate and hydroxyapatite. After the solution attains equilibrium at the pH, the hydroxyapatite is not deposited. In order to continue the deposition of hydroxyapatite it is necessary to maintain the pH of the solution at a weakly basic (for example pH 7.5-10).

Tetracalcium phosphate dissolves in distilled water to attain equilibrium as shown in the formula 3 (see Figure 7):



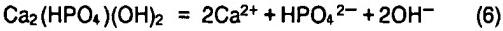
The equilibrated solution is a saturated aqueous solution of tetracalcium phosphate showing highly basic (pH of 12 or more), which is supersaturated with respect to hydroxyapatite.

35 Tetracalcium phosphate causes the following reaction as shown in the formula 4 to convert into hydroxyapatite in the equilibrated solution, and the formed hydroxyapatite is deposited:



40 When the reaction 4 proceeds, the pH of the solution rises because of the calcium hydroxide formation. In general, calcium phosphates are freely soluble in acidic solutions and slightly soluble in basic solutions. Therefore, since the rise of the pH of the solution results in decrease in the reaction rate, coagulation-hardening can not be expected.

45 Hydroxyapatite dissolves in distilled water by the reaction as shown in the formulae 5 and 6 to attain equilibrium:



In accordance with these reactions, the calcium hydroxide formed on the conversion of tetracalcium phosphate into hydroxyapatite is also able to produce hydroxyapatite via $\text{Ca}_2(\text{HPO}_4)(\text{OH})_2$ in the presence of secondary phosphate ion.

As can be seen in Figures 5 and 6, the pH of the neighborhood of the pastes kneaded with the disodium hydrogen phosphate-citric acid type buffer solutions rose gradually from pH about 6.5 for the paste using the tetracalcium phosphate powder and lay constantly at pH near 5.6 for that using the calcium hydrogen phosphate powder. However, for the TeDCPD pastes (Examples 31 and 32), the pH values were not there between the curves TeCP and DCPD throughout from the start of kneading and far from these

curves, showed less fluctuation, and lay constantly at a weakly basic (pH about 8.5).

In contrast, in the case of kneading with liquid components other than the disodium hydrogen phosphate-citric acid type buffer solutions, there were the following problems:

- As can be seen in Figure 7, the pH values of the neighborhood of the pastes kneaded with distilled water were stabilized at pH about 12.3 for the tetracalcium phosphate powder paste and at pH about 7.8 for the calcium hydrogen phosphate powder paste, and the TeDCPD paste (Reference Example 21) showed almost same behavior as the curve TeCP throughout from the start of kneading.

As can be seen in Figure 8, the pH values of the neighborhood of the pastes kneaded with the 20 mM aqueous orthophosphoric acid solution were stabilized at pH about 8 for the tetracalcium phosphate power paste, slightly fluctuated at near pH 5.5 for the calcium hydrogen phosphate powder paste, and lay constantly at pH about 8.3 throughout from the start of kneading for the TeDCPD paste (Reference Example 23).

As can be seen in Figure 9, the pH values of the neighborhood of the pastes kneaded with the 0.2 M aqueous disodium hydrogen phosphate solution rose from about 9.4 for the tetracalcium phosphate paste, lay constantly at pH about 8.6 for the calcium hydrogen phosphate paste, and were there between the curves TeCP and DCPD initially and showed the same behavior as the curve DCPD finally for the TeDCPD paste (Reference Example 24).

As can be seen in Figure 10, the pH values of the neighborhood of the pastes kneaded with the 0.4 M aqueous disodium hydrogen phosphate solution rose from pH about 9.2 for the tetracalcium phosphate paste, lay constantly at pH about 8.7 for the calcium hydrogen phosphate, and showed different behavior from the curves TeCP and DCPD for the TeDCPD paste (Reference Example 25). Thus, the pH values were once stabilized at pH about 9.5 initially, then lowered, and were stabilized at pH about 9 finally.

As can be seen in Figure 11, the pH values of the neighborhood of the pastes kneaded with the 0.1 M aqueous citric acid solution rose gradually from pH about 5.5 for the tetracalcium phosphate paste, slightly fluctuated at near pH 4 for the calcium hydrogen phosphate paste, and showed almost same behavior as the curve TeCP throughout from the start of kneading for the TeDCPD paste (Reference Example 26).

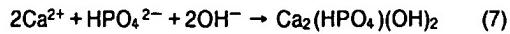
As can be seen in Figure 12, the pH values of the neighborhood of the pastes kneaded with the 0.2 M aqueous citric solution rose gradually from pH about 5 and 3.4 for the tetracalcium phosphate and calcium hydrogen phosphate pastes, respectively, and showed the same behavior as the curve TeCP for the TeDCPD paste (Reference Example 27).

The reason why the fluctuations of the pH of the neighborhood of the TeDCPD pastes are different between Figures 4 and 7-12 is because of the difference of the powder to liquid ratio used.

From the above results, when the kneading solutions are weakly acidic solutions, it is expected that tetracalcium phosphate in TeDCPD first dissolves in the solutions. Considering that when tetracalcium phosphate and calcium hydrogen phosphate were kneaded with the disodium hydrogen phosphate-citric acid type buffer solutions, respectively, the pH values of the neighborhood of these pastes were about 7 and 5.5, respectively, at 1 hour after the start of kneading, it is assumed that the dissolution of tetracalcium phosphate (formula 3) or the dissolution of tetracalcium phosphate and calcium hydrogen phosphate (formulae 1 and 3) was fast.

Since the dissolution shown the formulae 1 and 3 proceeds in an extremely short time because of the presence of the disodium hydrogen phosphate and citric acid, which are original solutes, and the elutes from tetracalcium phosphate and calcium hydrogen phosphate, equilibrium is reached immediately after kneading.

When the pH value of the solution is near 8.5 after equilibrium is reached, the reaction shown in the formula 2 proceeds until all of the calcium hydrogen phosphate convert into hydroxyapatite during the pH is maintained at near 8.5. The H_3PO_4 formed by the reaction comes to solution-equilibrium with the disodium hydrogen phosphate and citric acid during kneading. If the pH is still maintained at near 8.5, the reaction of the formula 4 also proceeds and the tetracalcium phosphate converts into hydroxyapatite. The $Ca(OH)_2$ produced in this reaction comes to solution-equilibrium with H_3PO_4 produced from calcium hydrogen phosphate and disodium hydrogen phosphate and citric acid in kneading solutions, and the reaction shown as the formulae 7 and 8 proceeds:



The hardening material described in said U.S. Patent No. 4518430 is a material in which calcium hydrogen phosphate and tetracalcium phosphate cause hydration-reaction by kneading the both phosphates

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with, for example, a diluted orthophosphoric acid solution to convert into hydroxyapatite, resulting in coagulation-hardening. This hydration-reaction occurs at pH about 7.5 at which the solubility curves of the both phosphates cross. In the hydration-reaction of calcium phosphates, the pH of the neighborhood of the pastes is largely fluctuated even when the condition such as mean particle size of powder, concentration of kneading solution, and powder to liquid ratio is slightly different. Because of this reason, it is difficult to maintain the pH of the neighborhood of the pastes at pH near 7.5 upon hydration-reaction of calcium hydrogen phosphate and tetracalcium phosphate. Even if the pH were maintained at the pH, the above condition would extremely be limited. As can be seen in Figures 4 and 8 showing the results of Reference Example 23, the difference of powder to liquid ratio provided a large difference in the pH of the neighborhood of the pastes even when the phosphoric acid solution of the same concentration was used.

In contrast, for the disodium hydrogen phosphate-citric acid type buffer solutions, the pH of the neighborhood of the pastes is not affected at all by the difference in the powder to liquid ratio and in the concentration of the solutes in the buffer solutions and is stabilized similarly.

As shown in Table 7, TeDCPD hardened in 6 and 8 minutes when kneaded with the disodium hydrogen phosphate solutions and did not harden when kneaded with the aqueous citric acid solutions. From these results, it is clear that, when TeDCPD is kneaded with disodium hydrogen phosphate-citric acid type buffer solutions to result in hardening, the necessary compound for hardening is the secondary phosphate ion. Therefore, the setting time can be controlled by regulating the pH and concentration of the solute in the kneading solution. This was also verified by the result that increase in concentration of secondary phosphate and organic acid ions caused faster setting time.

When kneading solution contains organic acid ions, a chelating reaction occurs between calcium and the ions in a small extent, and hence viscosity comes on kneading.

EXAMPLES 36-40

Hardening materials in the present invention are prepared using the combinations of the powder component obtained in the example 2 for preparation of powder component and the liquid components shown in Table 8.

REFERENCE EXAMPLES 28 and 29

Hardening materials are prepared using the combinations of the powder component in the example 2 and the liquid components shown in Table 8.

The materials in Examples 36-40 and Reference Examples 28 and 29 were kneaded using the powder to liquid ratio (powder weight/liquid weight) shown in Table 8 to examine kneading-processability, setting time, and pH of the neighborhood of the pastes (at 5 and 20 minutes after the start of kneading) by the above-described methods, and the adhesion to lentulo was also examined by the method described below. The results show in Table 8.

(Adhesion to lentulo)

The materials from which the pastes adhere uniformly to lentulo (a filling tool for dental root canal sealing materials) are judged as 'good adhesion'.

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TABLE 8

	EXAMPLE					REFERENCE EXAMPLE	
	36	37	38	39	40	28	29
Liquid Component	Soln. A	Soln. B	Soln. C	Soln. D	Soln. E	Soln. F	Soln. G
Powder/Liquid Wt. Ratio	1.2	1.2	1.2	1.2	1.2	1.2	1.6
Kneading-Processability	excellent	excellent	excellent	excellent	excellent	poor	poor
Adhesion	good	good	good	good	good	n.a.	n.a.
Setting Time (min.)	40	15	15	8	8	n.h.	n.h.
pH of Neighbor-	After 5 min.	8.56	8.57	8.33	8.34	8.47	13.15
hood of Paste	After 20 min.	8.70	8.67	8.47	8.11	8.16	10.30
						13.34	12.30
						11.00	12.51

Soln. A; buffer solution 1 in Table 5 containing 2.5 wt.% carboxymethyl cellulose sodium

Soln. B; buffer solution 2 in Table 5 containing 2.5 wt.% carboxymethyl cellulose sodium

Soln. C; buffer solution 2 in Table 5 containing 6.0 wt.% poly(sodium L-glutamate)

Soln. D; buffer solution 4 in Table 5 containing 2.5 wt.% carboxymethyl cellulose sodium

Soln. E; buffer solution 4 in Table 5 containing 6.0 wt.% poly(sodium L-glutamate)

Soln. F; distilled water containing 2.5 wt.% carboxymethyl cellulose sodium

Soln. G; 20 mM H₃PO₄ solution containing 2.5 wt.% carboxymethyl cellulose sodium

n.a.; not adhered to lentulo

n.h.; not hardened

Although, when TeDCPD was kneaded with the solutions containing secondary phosphate and organic acid ions, viscosity came slightly, the disodium hydrogen phosphate-citric acid type and disodium hydrogen phosphate-malic acid type buffer solutions containing carboxymethyl cellulose sodium or poly(sodium glutamate) were excellent as solutions providing cobwebbing property, as shown in Table 8. The addition of carboxymethyl cellulose sodium to distilled water or 20 mM orthophosphoric acid caused worse kneading-

processability and disappearance of hardenability.

In general, it is known that viscosity is affected strongly by pH value in case of using thickeners such as carboxymethyl cellulose sodium, poly(sodium glutamate) to enhance the liquid viscosity. Since such thickeners are effective to increase viscosity in near neutral regions, they show no effect when used for the 5 pastes showing high basic. In contrast, the addition of these thickeners to the secondary phosphate ion-organic acid ion type buffer solutions, which are the liquid components of the hardening materials wherein the pH of the neighborhood of their pastes stabilizes at weakly basic immediately after heating, is useful for enhancing viscosity and, in addition, for providing pastes with the cobwebbing property.

10 Claims

1. A hardenable, biocompatible composition for repairing living hard tissue comprising a powder component containing $\text{Ca}_4\text{P}_2\text{O}_9$ and at least one of CaHPO_4 and $\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$ at a molar ratio of Ca:P of 1.16- 1.95; and
- 15 a liquid component containing a secondary phosphate ion in a concentration of 1mM to 2M as a secondary phosphate and an organic acid ion in a concentration of 1mM to 2M as an organic acid or as a total of an organic acid and its salt, wherein a weight ratio of the powder component : the liquid component is 0.5-4.0.
- 20 2. A hardenable, biocompatible composition as claimed in Claim 1, having a pH range of about 8-9.
3. A hardenable, biocompatible composition as claimed in Claim 1 or Claim 2, wherein the secondary phosphate ion source is disodium hydrogen phosphate, dipotassium hydrogen phosphate, or diammonium hydrogen phosphate.
- 25 4. A hardenable, biocompatible composition as claimed in any preceding Claim, wherein the powder component has a mean particle size range of about 0.1-50 μm .
- 30 5. A hardenable, biocompatible composition as defined in any preceding Claim, wherein the organic acid is citric acid, malic acid, malonic acid, maleic acid, fumaric acid, fumaleic acid, lactic acid, acetic acid, oxalacetic acid, isocitric acid, aconitic acid, succinic acid, acrylic acid homopolymer, acrylic acid-itaconic acid copolymer, or salts thereof.
- 35 6. A hardenable, biocompatible composition as claimed in any of Claims 1-4, wherein the organic acid is citric acid, metabolism products of citric acid, or salts thereof.
7. A hardenable, biocompatible composition as claimed in any preceding Claim, further comprising at 40 least one thickener selected from the group consisting of carboxy methyl cellulose, sodium carboxymethyl cellulose, hydroxypropyl cellulose, polyvinyl alcohol, polyglutamic acid, polyglutamic acid salt, polyaspartic acid and polyaspartic acid salt, wherein the concentration of polyglutamic acid and polyglutamic acid salt is 0.5-15% by weight and the concentration of the other thickeners is 0.5-10% by weight.
- 45 8. A hardenable, biocompatible composition as claimed in any of Claims 1-6 further comprising at least one thickener selected from the group consisting of sodium carboxymethyl cellulose, hydroxypropyl cellulose, polyglutamic acid, and polyglutamic acid salt.
9. A hardenable, biocompatible composition as claimed in any preceding Claim, further comprising an X-ray contrast medium.

50 Patentansprüche

1. Härtbare, biokompatible Zusammensetzung zum Ausbessern von lebendem Hartgewebe, bestehend aus einer Pulverkomponente, die $\text{Ca}_4\text{P}_2\text{O}_9$ und wenigstens eines von CaHPO_4 und $\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$ mit einem molaren Verhältnis von Ca:P von 1.16 bis 1.95 enthält; und
- 55 einer flüssigen Komponente, die ein sekundäres Phosphat mit einer Konzentration von 1mM bis 2M als ein sekundäres Phosphat und ein organisches Säureion mit einer Konzentration von 1mM bis 2M

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als eine organische Säure oder als eine Gesamtheit einer organischen Säure und ihres Salzes enthält, wobei ein Gewichtsverhältnis der Pulverkomponente zu der flüssigen Komponente 0.5 bis 4.0 beträgt.

2. Härtbare, biokompatible Zusammensetzung nach Anspruch 1, die einen pH Bereich von etwa 8 bis 9 aufweist.
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3. Härtbare, biokompatible Zusammensetzung nach Anspruch 1 oder Anspruch 2, bei welcher die Quelle des sekundären Phosphats Dinatriumhydrogenphosphat, Dikaliumhydrogenphosphat oder Diammoniumhydrogenphosphat ist.
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4. Härtbare, biokompatible Zusammensetzung nach einem der vorhergehenden Ansprüche, bei welcher die Pulverkomponente eine mittlere Teilchengröße im Bereich von etwa 0.1 bis 50 µm hat.
15
5. Härtbare, biokompatible Zusammensetzung nach einem der vorhergehenden Ansprüche, bei welcher die organische Säure Zitronensäure, Apfelsäure, Malonsäure, Aleinsäure, Fumarsäure, Fumaleinsäure, Milchsäure, Essigsäure, Oxalesigsäure, Isozitronensäure, Aconitsäure, Succinsäure, Acrylsäure-Homopolymer, Acrylsäure-Itaconsäure-Copolymer oder deren Salze ist bzw. sind.
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6. Härtbare, biokompatible Zusammensetzung nach einem der Ansprüche 1 bis 4, bei welcher die organische Säure Zitronensäure, Stoffwechsel-Produkte der Zitronensäure oder deren Salze ist bzw. sind.
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7. Härtbare, biokompatible Zusammensetzung nach einem der vorhergehenden Ansprüche, welche weiterhin wenigstens ein Verdickungsmittel aufweist, ausgewählt aus der Gruppe, bestehend aus Carboxymethylzellulose, Natriumcarboxymethylzellulose, Hydroxypropylzellulose, Polyvinylalkohol, Polyglutaminsäure, Polyglutaminsäuresalz, Polyasparaginsäure und Polyasparaginsäuresalz, wobei die Konzentration der Polyglutaminsäure und des Polyglutaminsäuresalzes 0.5 - 15 Gew.-% und die Konzentration der anderen Verdickungsmittel 0.5 - 10 Gew.-% beträgt.
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8. Härtbare, biokompatible Zusammensetzung nach einem der Ansprüche 1 bis 6, welche wenigstens ein Verdickungsmittel enthält, ausgewählt von der Gruppe, bestehend aus Natriumcarboxymethylzellulose, Hydroxypropylzellulose, Polyglutaminsäure und Polyglutaminsäuresalz.
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9. Härtbare, biokompatible Zusammensetzung nach einem der vorhergehenden Ansprüche, welche weiterhin ein Röntgenstrahl-Kontrastmittel enthält.

Revendications

1. Composition durcissable, biocompatible pour réparer des tissus vivants durs, comprenant un composant en poudre contenant Ca₄P₂O₉ et au moins l'un de CaHPO₄ et CaHPO₄.2H₂O à un taux molaire de Ca:P de 1,16- 1,95; et un composant liquide contenant un ion phosphate secondaire dans une concentration de 1mM à 2M en tant que phosphate secondaire et un ion d'acide organique dans une concentration de 1mM à 2M en tant qu'acide organique ou en tant que total entre un acide organique et son sel, dans laquelle la proportion en poids du composant en poudre par rapport au composant liquide est de 0,5 à 4,0.
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2. Composition durcissable, biocompatible telle que revendiquée dans la revendication 1, ayant une plage de pH d'environ 8-9.
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3. Composition durcissable, biocompatible telle que revendiquée dans la revendication 1 ou la revendication 2, dans laquelle la source d'ion phosphate secondaire est le phosphate d'hydrogène disodique, le phosphate d'hydrogène dipotassique, ou le phosphate d'hydrogène diammonium.
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4. Composition durcissable, biocompatible telle que revendiquée dans l'une quelconque des revendications précédentes, dans laquelle le composant en poudre a une plage de taille de particule principale d'environ 0,1-50µm.
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5. Composition durcissable, biocompatible telle que définie dans l'une quelconque des revendications précédentes, dans laquelle l'acide organique est l'acide citrique, l'acide malique, l'acide malonique, l'acide maléique, l'acide fumarique, l'acide fumaléique, l'acide lactique, l'acide acétique, l'acide oxalacétique, l'acide isocitrique, l'acide aconitrique, l'acide succinique, l'homopolymère d'acide acrylique, un copolymère d'acide itaconique, ou leurs sels.
10. Composition durcissable, biocompatible telle que revendiquée dans l'une quelconque des revendications 1 à 4, dans laquelle l'acide organique est l'acide citrique, des produits de métabolisme de l'acide citrique, ou leurs sels.
15. Composition durcissable, biocompatible telle que revendiquée dans l'une quelconque des revendications précédentes, comprenant en outre au moins un épaississant sélectionné à partir du groupe consistant dans la carboxyméthylcellulose, la carboxyméthylcellulose de sodium, l'hydroxypropylcellulose, l'alcool polyvinyle, l'acide polyglutamique, un sel d'acide polyglutamique, l'acide polyaspartique et un sel de l'acide polyaspartique, dans laquelle la concentration d'acide polyglutamique et de sel d'acide polyglutamique est de 0,5 à 15 % en poids et la concentration des autres épaississants est de 0,5 à 10 % en poids.
20. Composition durcissable, biocompatible telle que revendiquée dans l'une quelconque des revendications 1 à 6, comprenant en outre au moins un épaississant sélectionné à partir du groupe consistant dans la carboxyméthylcellulose de sodium, l'hydroxypropylcellulose, l'acide polyglutamique, et un sel de l'acide polyglutamique.
25. Composition durcissable, biocompatible telle que revendiquée dans l'une quelconque des revendications précédentes, comprenant en outre un moyen de contraste aux rayons X.

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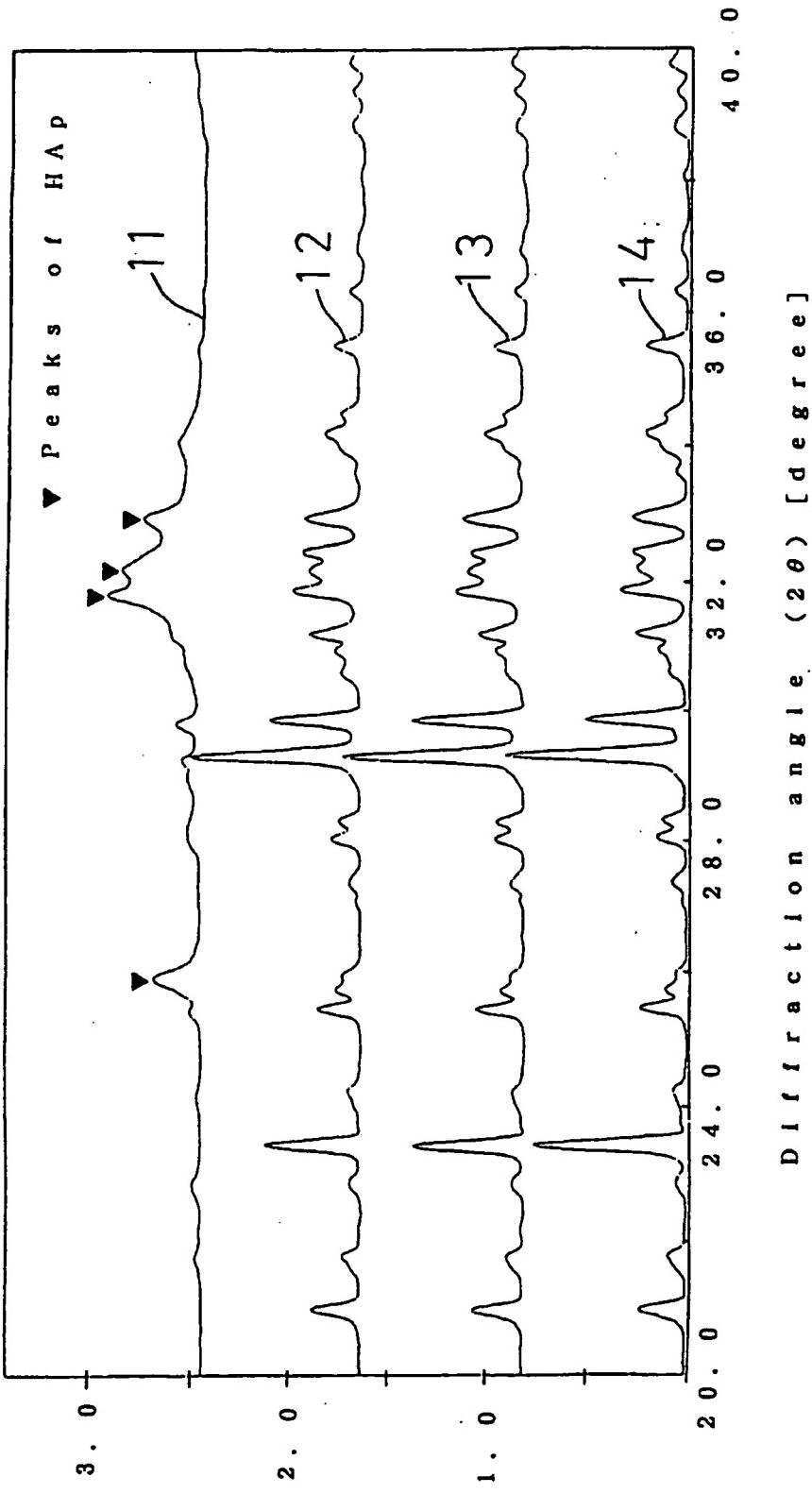
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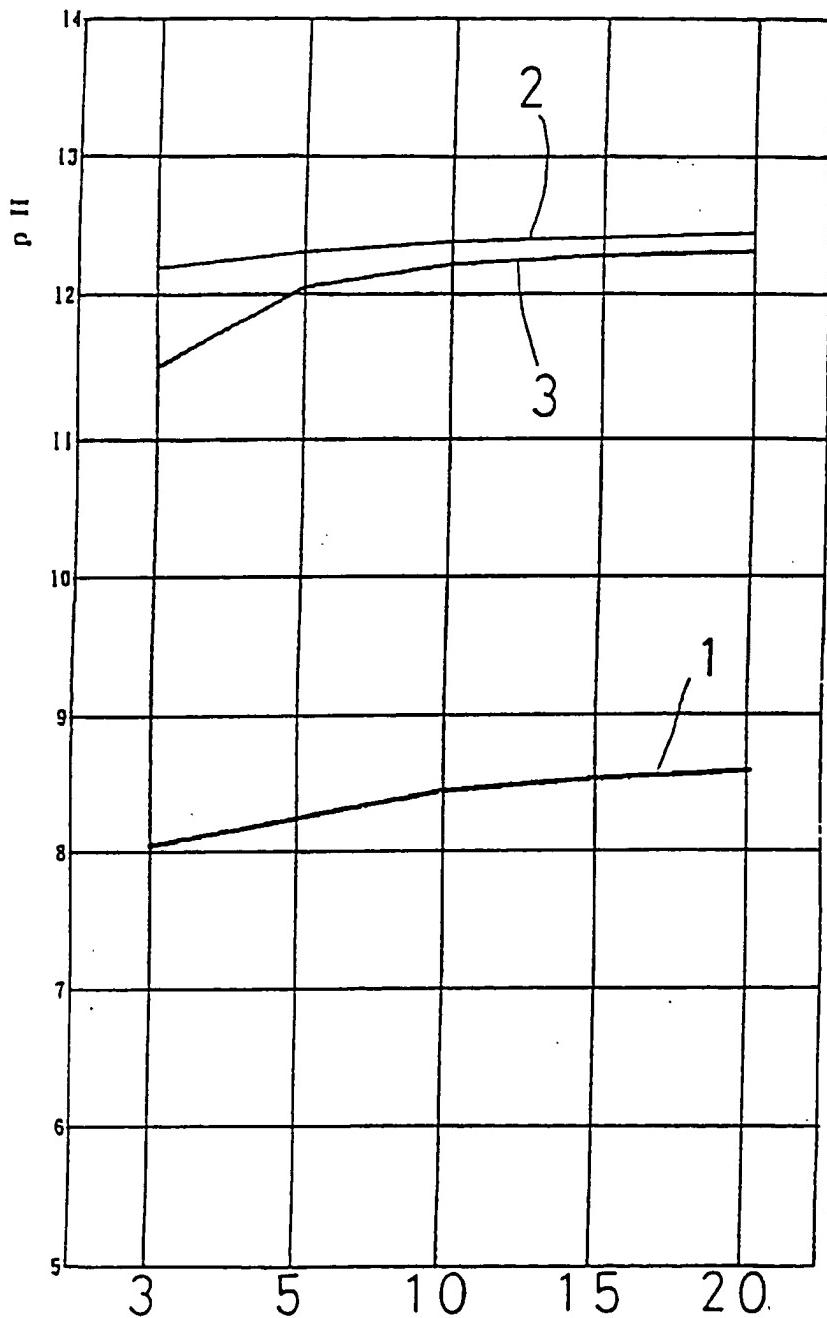
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Fig. 1



D I F F R A C T I O N I n t e n s i t y I [K (c p s)]

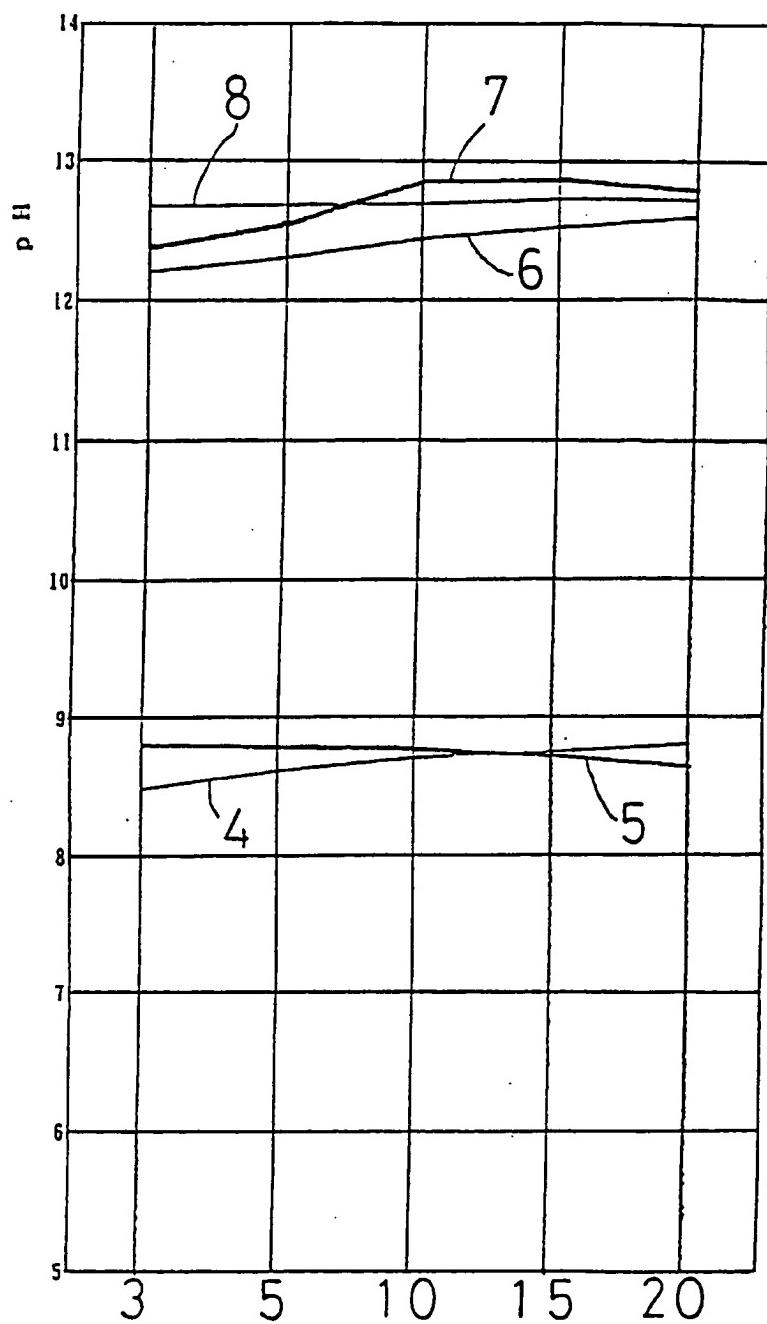
F i g . 2



T i m e f r o m t h e s t a r t o f k n e a d i n g .

(m i n) .

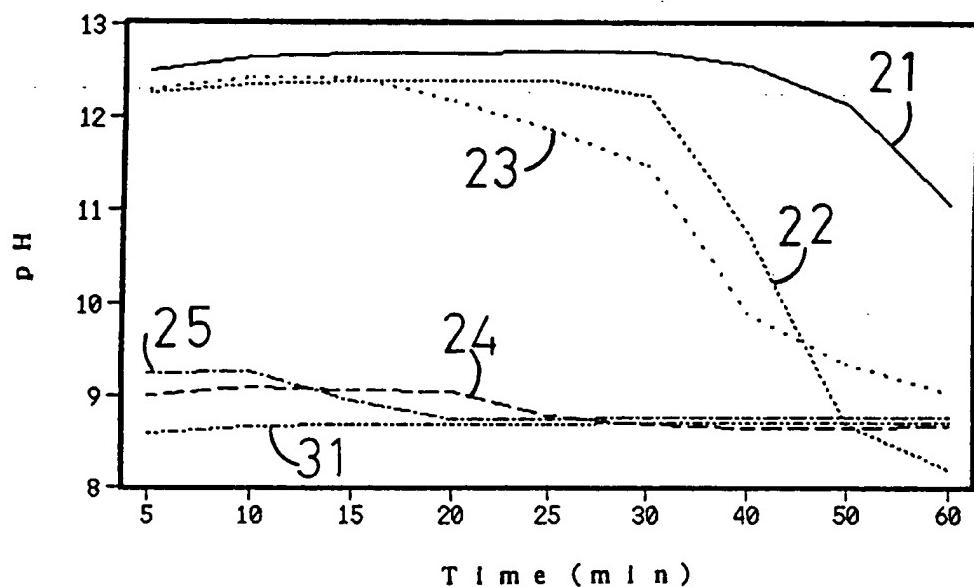
F i g . 3



T i m e f r o m t h e s t a r t o f k n e a d i n g
(m i n)

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F i g . 4



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Fig. 5

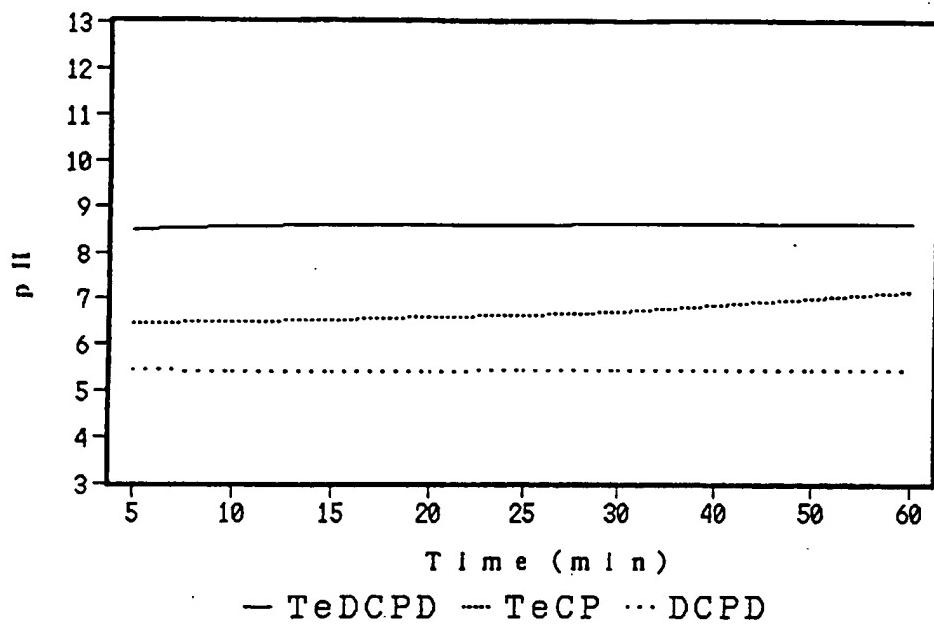
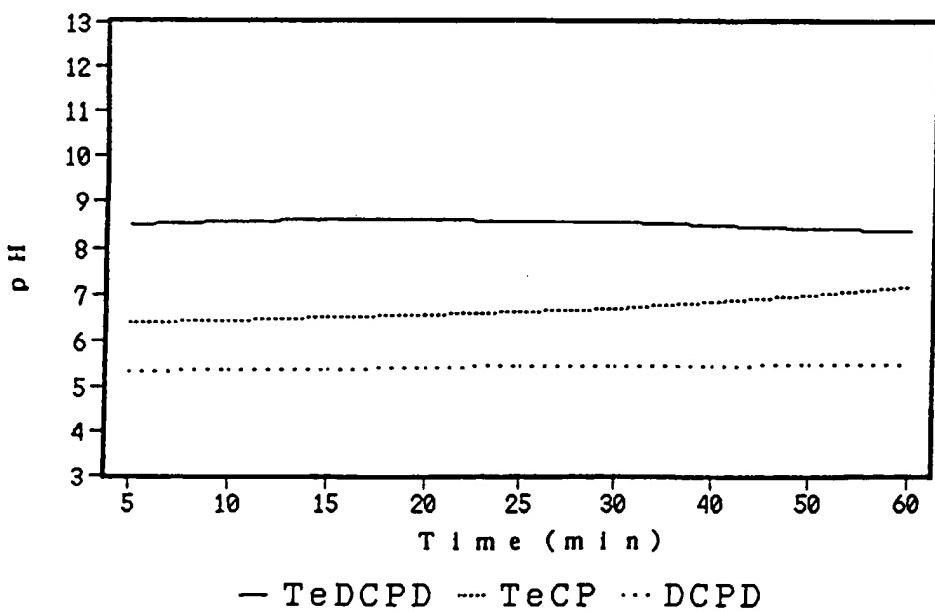


Fig. 6



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Fig. 7

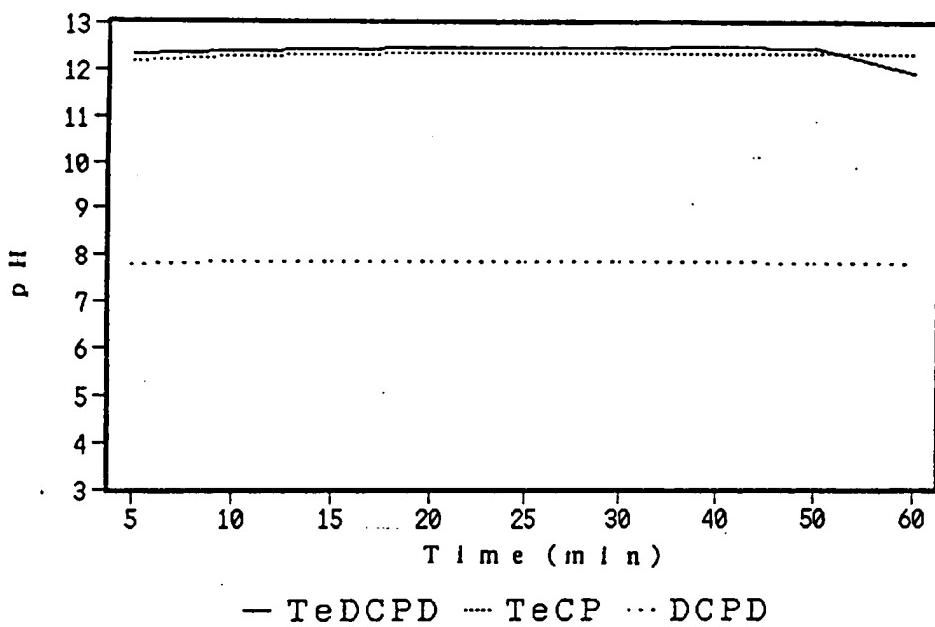
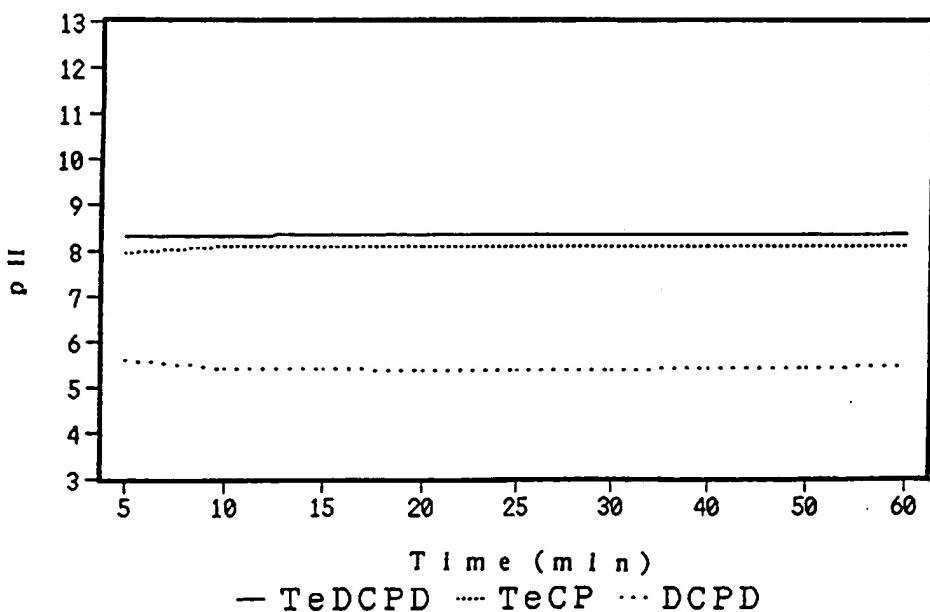


Fig. 8



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Fig. 9

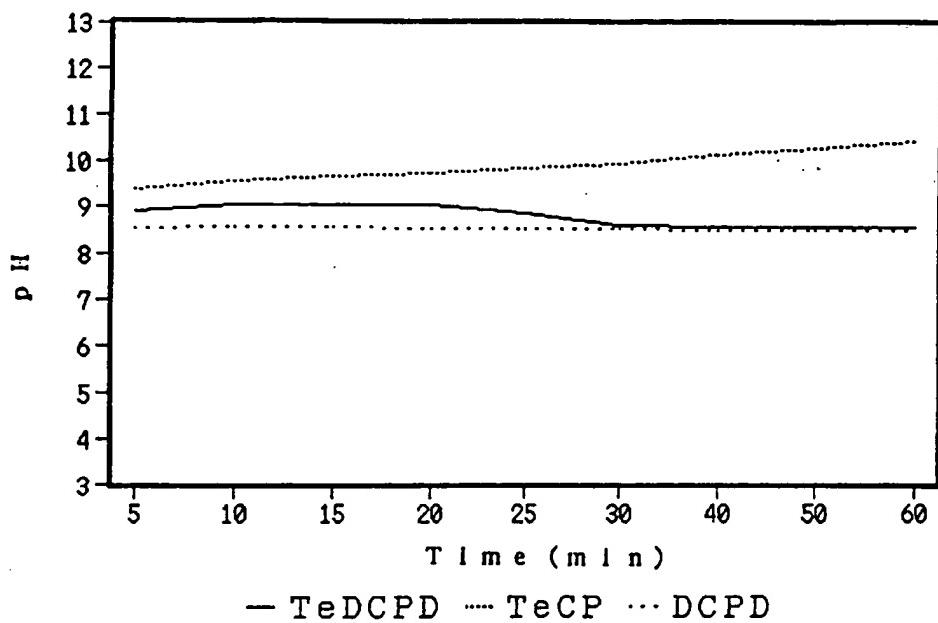
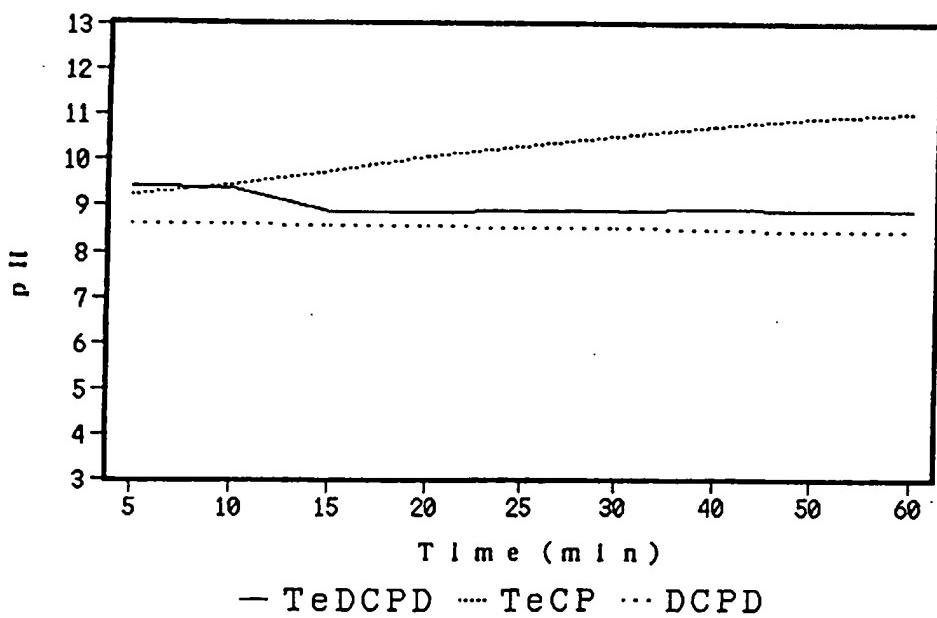


Fig. 10



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Fig. 11

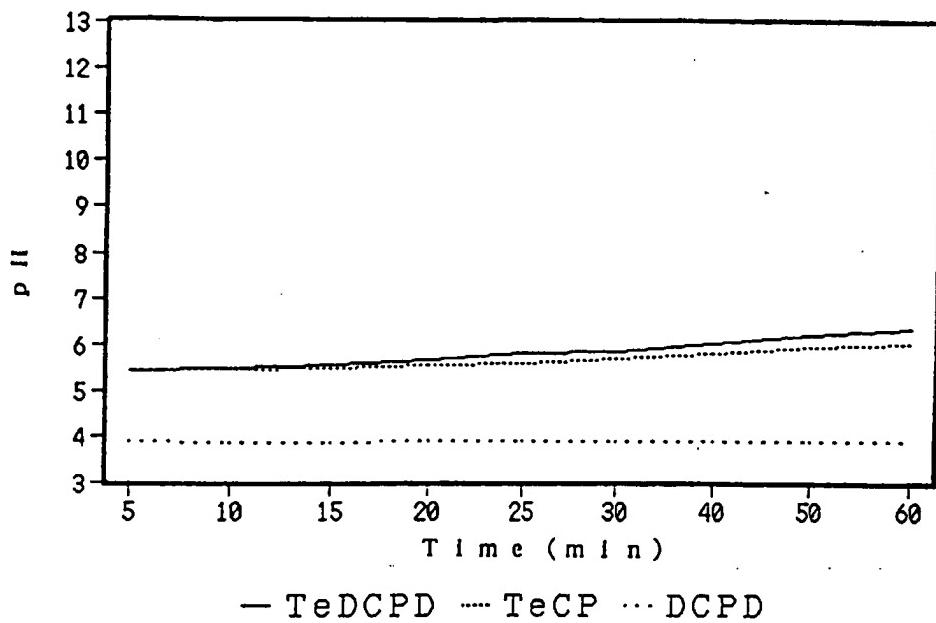


Fig. 12

